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Form PTO-1390 (REV 10-95)		U. S. Department of Commerce Patent and Trademark Office	ATTORNEY'S DOCKET NUMBER <b>GT/3-21914/A/AC 524/PCT</b>
<b>TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371</b>			U.S. APPLICATION NO. (If known, see 37 CFR 1.5)
			<b>10/030535</b>
INTERNATIONAL APPLICATION NO. <b>EP 00/06292</b>	INTERNATIONAL FILING DATE <b>July 5, 2000</b>	PRIORITY DATE CLAIMED <b>July 19, 1999</b>	
TITLE OF INVENTION <b>PROCESS FOR FLOCCULATING SUSPENSIONS</b>			
APPLICANT(S) FOR DO/EO/US <b>Steven Weir and Mark John Stringer</b>			

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☐ This express request to begin national examination procedures (35 U.S.C. 371(f) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39 (1).
4. ☒ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
  - a. ☐ is transmitted herewith (required only if not transmitted by the International Bureau).
  - b. ☒ has been transmitted by the International Bureau. (See attached Form PCT/IB/308)
  - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☐ A translation of the International Application into English 35 U.S.C. 371(c)(2)).
7. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3)).
  - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
  - b. ☐ have been transmitted by the International Bureau.
  - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
  - d. ☒ have not been made and will not be made.
8. ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371 (c)(3)).
9. ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11. to 16. below concern document(s) or information included.

11. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☒ **A FIRST preliminary amendment.**  
☐ A SECOND or SUBSEQUENT preliminary amendment.
14. ☐ A substitute specification.
15. ☐ A change of power of attorney and/or address letter.
16. ☒ Other items or information: (See attached Form PCT/ISA/210)

U.S. APPLICATION NO. (if known, see 37 CFR 1.55) <b>10/030535</b>		INTERNATIONAL APPLICATION NO. <b>EP 00106292</b>		ATTORNEY'S DOCKET NUMBER <b>GT/3-21914/A/AC 524/PCT</b>	
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17. <input checked="" type="checkbox"/> The following fees are submitted:				<b>CALCULATIONS PTO USE ONLY</b>	
<b>BASIC NATIONAL FEE (37 CFR 1.492(a) (1)-(5)):</b>					
Search Report has been prepared by the EPO or JPO .....				\$890.00	
International preliminary examination fee paid to USPTO (37 CFR 1.482) .....				\$890.00	
.....				\$710.00	
No international preliminary examination fee paid to USPTO (37 CFR 1.482) but international search fee paid to USPTO (37 CFR 1.445(a)(2)). .....				\$740.00	
Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO. ....				\$1040.00	
International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(2)-(4). ....				\$100.00	
<b>ENTER APPROPRIATE BASIC FEE AMOUNT =</b>				\$890.00	
Surcharge of \$130.00 for furnishing the oath of declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(e)).				\$	
CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE		
Total claims	19 - 20 =	0	X \$18.00	\$	
Independent claims	1 - 3 =	0	X \$84.00	\$	
MULTIPLE DEPENDENT CLAIM(S) (if applicable)			+ \$280.00	\$	
<b>TOTAL OF ABOVE CALCULATIONS =</b>				\$890.00	
Reduction of 1/2 for filing by small entity, if applicable. Verified Small Entity Statement must also be filed (Note 37 CFR 1.9, 1.27, 1.28).				\$	
<b>SUBTOTAL =</b>				\$890.00	
Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)).				\$	
<b>TOTAL NATIONAL FEE =</b>				\$890.00	
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property				\$	
<b>TOTAL FEES ENCLOSED =</b>				\$	
				Amount to be:	\$
				refunded	
				charged	\$890.00


a. ☒ Please charge my Deposit Account No. 03-1935 in the amount of \$890.00 to cover the above fees. A duplicate copy of this sheet is enclosed.

b. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 03-1935. A duplicate copy of this sheet is enclosed.

**NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.**

PLEASE ASSOCIATE THE ATTACHED APPLICATION WITH **CUSTOMER NUMBER 000324** AND SEND ALL CORRESPONDENCE TO:

JoAnn Villamizar  
Patent Department  
540 White Plains Road  
P.O. Box 2005  
Tarrytown, NY 10591-9005

  
 SIGNATURE  
  
 David R. Crichton  
 NAME  
  
 37,300  
 REGISTRATION NUMBER

CASE GT/3/21914/A/AC 524

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

IN RE APPLICATION OF

Group Art Unit:

STEVEN WEIR ET AL.

Examiner:

APPLICATION NO: Not Yet Assigned

FILED: Concurrently Herewith

FOR: PROCESS FOR FLOCCULATING

SUSPENSIONS

Assistant Commissioner for Patents

Washington, D.C. 20231

PRELIMINARY AMENDMENT

Dear Sir:

Applicants present the instant Preliminary Amendment for entry and consideration in order to place the instant application in better condition for examination on its merits and for allowance.

The Commissioner is authorized to charge any fee due, or credit any overcharge, as a result of this Preliminary Amendment to Deposit Account No. 03-1935.

Please amend the above-identified patent application, without prejudice, as follows:

IN THE CLAIMS:

**Replace claims 5-14, 16 and 19 by replacement as follows:**

5. A process according to claim 1 in which the dilute aqueous solution has a concentration of polymer of below 0.3% by weight.
6. A process according to claim 1 in which the dilute aqueous polymer solution comprises a cationic polymer, an anionic polymer or a nonionic polymer.
7. A process according to claim 1 in which the concentrated aqueous solution has a concentration of polymer of between 0.4 and 1.0%, by weight.
8. A process according to claim 1 in which the concentrated aqueous solution comprises a cationic polymer, an anionic polymer or a nonionic polymer.
9. A process according to claim 1 in which the polymer dissolved in the concentrated solution is either co-ionic with the polymer dissolved in the dilute solution or non-ionic.
10. A process according to claim 1 in which polymer dissolved in either the dilute solution or the concentrated solution is cationic and has been formed from a monomer or blend of monomers comprising at least one cationic monomer selected from the group consisting of quaternary ammonium and acid salts of dimethylaminoethyl (meth) acrylate, quaternary ammonium and acid salts of dimethylaminoethyl (meth) acrylamide and diallyldimethyl ammonium chloride.
11. A process according to claim 1 in which the polymer(s) dissolved in either the dilute solution or the concentrated solution is anionic and has been formed from a monomer or blend of monomers comprising at least one anionic monomer selected from the group consisting of (meth) acrylic acid, 2-acrylamido-2-methylpropane sulphonic acid, alkali metal and ammonium salts thereof.

12. A process according to claim 1 in which the polymer(s) dissolved in either the dilute solution or the concentrated solution is nonionic and has been formed from acrylamide or methacrylamide.

13. A process according to claim 1 in which the cationic polymer dissolved in each of the dilute and concentrated aqueous solutions is a copolymer of acrylamide and at least one cationic monomer selected from the group consisting of quaternary ammonium and acid salts of dimethylaminoethyl (meth) acrylate, quaternary ammonium and acid salts of dimethylaminoethyl (meth) acrylamide and diallyldimethyl ammonium chloride, having an intrinsic viscosity of at least 4 dl/g.

14. A process according to claim 2 in which the aqueous composition comprising the dilute aqueous solution of polymer and the concentrated solution of polymer is formed by introducing the concentrated solution of polymer into a flowing stream of the dilute aqueous solution of polymer.

16. A process according to claim 2 in which the aqueous composition is formed by,  
 (a) passing a concentrated solution of polymer to a dilution where the solution is combined with dilution water to form a dilute solution,  
 (b) passing the diluted solution through a mixing stage, selected from pumping and screening stages, and  
 (c) introducing a concentrated solution of polymer into the dilute aqueous solution.

19. A process according to claim 1 in which the dewatering process is selected from the group consisting of dewatering sewage sludge, dewatering a mineral suspension, dewatering a paper mill sludge, dewatering a deinked cellulosic sludge and a papermaking process.

Remarks

Upon entry of the instant Preliminary Amendment, claims 1-19 are pending. Multiple dependencies have been eliminated from the claims without prejudice to the filing of claims directed thereto in the instant or a subsequent application. The amendment is strictly a matter of form. No new matter has been added.

In view of the foregoing amendments, Applicants aver that the instant claims are now in better condition for examination on the merits and for allowance. Early favorable action is respectfully solicited. If minor amendments will further prosecution, Applicants request that the Examiner contact the undersigned representative.

Respectfully submitted



David R. Crichton  
Attorney for Applicants  
Reg. No. 37,300

Ciba Specialty Chemicals Corporation  
540 White Plains Road  
P.O. Box 2005  
Tarrytown, New York 10591-9005  
Tel: 914-785-7124  
Fax: 914-785-7102  
DRC/

### Amended Claims with Mark-Ups

5. A process according to ~~any of claims 1 to 4~~claim 1 in which the dilute aqueous solution has a concentration of polymer of below 0.3% by weight.
6. A process according to ~~any one of claims 1 to 5~~claim 1 in which the dilute aqueous polymer solution comprises a cationic polymer, an anionic polymer or a nonionic polymer.
7. A process according to ~~any of claims 1 to 6~~claim 1 in which the concentrated aqueous solution has a concentration of polymer of between 0.4 and 1.0%, by weight.
8. A process according to ~~any one of claims 1 to 7~~claim 1 in which the concentrated aqueous solution comprises a cationic polymer, an anionic polymer or a nonionic polymer.
9. A process according to ~~any one of claims 1 to 8~~claim 1 in which the polymer dissolved in the concentrated solution is either co-ionic with the polymer dissolved in the dilute solution or non-ionic.
10. A process according to ~~any one of claims 1 to 9~~claim 1 in which polymer dissolved in either the dilute solution or the concentrated solution is cationic and has been formed from a monomer or blend of monomers comprising at least one cationic monomer selected from the group consisting of quaternary ammonium and acid salts of dimethylaminoethyl (meth) acrylate, quaternary ammonium and acid salts of dimethylaminoethyl (meth) acrylamide and diallyldimethyl ammonium chloride.
11. A process according to ~~any one of claims 1 to 10~~claim 1 in which the polymer(s) dissolved in either the dilute solution or the concentrated solution is anionic and has been formed from a monomer or blend of monomers comprising at least one anionic monomer selected from the group consisting of (meth) acrylic acid, 2-acrylamido-2-methylpropane sulphonic acid, alkali metal and ammonium salts thereof.

12. A process according to ~~any one of claims 1 to 11~~claim 1 in which the polymer(s) dissolved in either the dilute solution or the concentrated solution is nonionic and has been formed from acrylamide or methacrylamide.

13. A process according to ~~any one of claims 1 to 12~~claim 1 in which the cationic polymer dissolved in each of the dilute and concentrated aqueous solutions is a copolymer of acrylamide and at least one cationic monomer selected from the group consisting of quaternary ammonium and acid salts of dimethylaminoethyl (meth) acrylate, quaternary ammonium and acid salts of dimethylaminoethyl (meth) acrylamide and diallyldimethyl ammonium chloride, having an intrinsic viscosity of at least 4 dl/g.

14. A process according to ~~any one of claims 2 to 13~~claim 2 in which the aqueous composition comprising the dilute aqueous solution of polymer and the concentrated solution of polymer is formed by introducing the concentrated solution of polymer into a flowing stream of the dilute aqueous solution of polymer.

16. A process according to ~~any one of claims 2 to 15~~claim 2 in which the aqueous composition is formed by,

- (a) passing a concentrated solution of polymer to a dilution where the solution is combined with dilution water to form a dilute solution,
- (b) passing the diluted solution through a mixing stage, selected from pumping and screening stages, and
- (c) introducing a concentrated solution of polymer into the dilute aqueous solution.

19. A process according to ~~any one of claims 1 to 18~~claim 1 in which the dewatering process is selected from the group consisting of dewatering sewage sludge, dewatering a mineral suspension, dewatering a paper mill sludge, dewatering a deinked cellulosic sludge and a papermaking process.



### Process For Flocculating Suspensions

This invention relates to processes of flocculating aqueous suspensions in order to effect separation of solids from said suspension.

It is well known to apply polymeric flocculants to aqueous suspensions in order to separate solids from the suspension. For instance it is common practice to flocculate and then dewater suspensions containing either suspended solid organic material or mineral solids. For instance it is common practice to flocculate sludges such as sewage sludge, waste waters, textile industry effluents, red mud from the Bayer Alumina process and suspensions of coal tailings etc. Flocculants are also commonly used in paper-making processes by addition of polymeric flocculants to the cellulosic suspension. Flocculation is usually achieved by mixing into the suspension polymeric flocculant, allowing the suspended particles to flocculate and then dewatering the flocculated suspension. In papermaking this removal of water from the cellulosic suspension is often referred to as draining.

High molecular weight polymeric flocculants are commonly used for this purpose. High molecular weight flocculants may be cationic, anionic, nonionic or amphoteric in nature. The choice of polymeric flocculant will largely depend upon the substrate, which is being treated. For instance it is common practice to use high molecular weight cationic flocculants to treat aqueous suspensions comprising suspended organic material, for instance sewage sludge. In paper-making it is known to use either cationic, nonionic, anionic or amphoteric flocculants. Flocculation of mineral suspensions is frequently effected by use of anionic flocculants.

It is also known to use two different polymeric flocculants in the same process. The flocculants may have the same charge (co-ionic). For instance in commercial practice in the dewatering sewage sludge these may be co-ionic. In other processes it is known to apply two polymers of opposite charge (counter-ionic). Where two polymeric flocculants are applied to an aqueous suspension they may be added to simultaneously or more usually sequentially.

It is standard practice to apply polymers as aqueous solutions to flocculate suspensions containing suspended organic material. Generally the solutions of polymers are relatively dilute, for instance below 0.5%, often below 0.3% and usually 0.1% to below 0.2% by weight.

Polymers are usually provided as a solid particulate product or as a reverse phase dispersion or emulsion. It is standard practice to dissolve the polymer into water by dispersing the polymer particles in a flowing stream of water in the case of the solid particulate product or in the case of the emulsion or dispersion, inversion into water, by use of activator surfactants. The polymer solution thus formed is frequently at a concentration above 0.3%, often in the range 0.4% to 1% and usually about 0.5%. This more concentrated solution of polymer may be too concentrated to add directly to the suspension in many instances, since received wisdom suggests that there would be inadequate distribution of the flocculant throughout the suspension and as a consequence the flocculation process would be impaired.

It is therefore common practice to first of all provide a more concentrated solution of polymer and then dilute the polymer solution prior to application. Often the diluted solution will have a concentration of below 0.2%, for instance within the range 0.05 to 0.2% by weight and frequently between 0.1 and 0.2% by weight. This dilute solution of polymer is normally metered directly into the suspension prior to the dewatering stage.

There is a desire to improve the efficiency of the flocculation processes, to either bring about an increased dewatering effect, such as higher cake solids or in the alternative achieve a constant acceptable level of dewatering efficiency but using a lower dose of flocculant. This is true in a variety of flocculation processes, including dewatering of sewage sludge, slurries of coal tailings, red mud and in papermaking.

It would therefore be desirable to provide an improved method of flocculating and dewatering aqueous suspensions of solids, in particular to provide increased dryness of the dewatered solids for an equivalent dose of flocculant or to provide the same degree of dryness of dewatered solids but using a reduced dose of flocculant. It would also be desirable to provide a process which provides faster dewatering.

The invention relates to a process of flocculating and dewatering an aqueous suspension of suspended solids comprising, introducing into the suspension,

- (a) a concentrated polymer solution and,
- (b) a dilute polymer solution,

characterised in that the concentrated and dilute polymer solutions are introduced into the substrate substantially simultaneously.

The concentrated and dilute solutions may be metered directly into the suspension as separate solutions. By substantially simultaneously the two solutions should be added at approximately the same dosing point. Where the concentrated and dilute solutions are added to the suspension separately, they may be added in either order. For instance if the dilute solution is added first the concentrated polymer may be added after flocculation has commenced but should be added before the dewatering stage and before any high shear stage, such as pumping or screening stages. Alternatively, it may be desirable to add the dilute polymer solution after the addition of the concentrated polymer solution. When the dilute and concentrated polymer solutions are added separately it may be appropriate to allow or apply some degree mixing between the dosing stages in order to allow the first polymer dose to become distributed throughout the suspension solids. This mixing may for instance include allowing the treated suspension to pass some distance along a flow line which optionally contains bends, baffles, constrictions or other features which induce gentle mixing.

Preferably the concentrated and dilute polymer solutions are introduced simultaneously.

More preferably the concentrated and dilute polymer solutions are introduced into the suspension as an aqueous composition comprising a dilute aqueous solution of polymer and a concentrated solution of polymer. The aqueous composition should comprise both the dilute and the concentrated polymer solutions as discrete components. Thus it is desired that the dilute solution and concentrated solution exist as substantially discrete components of the aqueous composition.

The aqueous composition preferably comprises the dilute aqueous solution of polymer in an amount of from 20 to 99%, based on weight of polymer, and the concentrated polymer solution in an amount of from 1 to 80%, based on weight of polymer. For some applications, such as for rotary vacuum filtration coal tailings slurries it may be appropriate to use a ratio of concentrated solution to dilute polymer solution of around 75:25. However in most other applications the ratio of concentrated polymer solution to dilute polymer solution would generally be in the range 1:99 to 40:60.

The aqueous composition comprising concentrated and dilute solutions may be of any significantly different concentrations provided that the respective concentrations are not substantially the same such that the two solutions would immediately form a homogenous single solution. Preferably the concentrated solution should be at least twice the concentration of the diluted solution. More preferably the concentrated solution should be at least 4 or 5 times the concentration of the dilute aqueous solution.

The dilute aqueous solution of polymer desirably has a concentration of polymer of below 0.5%, preferably below 0.3% by weight. More preferably the concentration of the dilute solution is in the range 0.05 to 0.2%, most preferably around 0.1% by weight.

According to the invention the polymer dissolved in the dilute aqueous polymer solution may be either cationic, anionic or non-ionic.

The concentrated aqueous solution component according to the invention desirably has a concentration of polymer above 0.3% by weight, preferably between 0.4 and 1.0% by weight. More preferably the concentration of the concentrated solution is in the range 0.5 to 1.0%. According to the invention the polymer dissolved in the concentrated aqueous polymer solution may be either cationic, anionic or non-ionic. The polymer dissolved in the concentrated polymer solution is preferably either co-ionic with the polymer dissolved in the dilute solution or non-ionic. In another preferred form the polymer dissolved in the dilute solution is non-ionic and the polymer dissolved in the concentrated polymer solution is cationic, anionic or non-ionic.

When the polymer dissolved in either the dilute solution or concentrated solution is cationic, said cationic polymer may be formed by polymerisation of at least one cationic monomer alone or with other monomers. Suitable cationic monomers include quaternary ammonium or acid salts of monomers which contain amine groups. Preferably the cationic polymer is formed from a monomer or blend of monomers comprising at least one cationic monomer selected from the group consisting of quaternary ammonium and acid salts of dimethylaminoethyl (meth) acrylate, quaternary ammonium and acid salts of dimethylaminoethyl (meth) acrylamide and diallyldimethyl ammonium chloride. The cationic monomers may be homopolymerised or copolymerised with other monomers, for instance acrylamide. In addition to vinyl addition polymers, the cationic polymer may include polymers obtained by condensation or addition reactions. For instance suitable cationic polymers include adducts of amines with epihalohydrins or dihaloalkanes, polyamides and polyethylene imines.

In the case where the polymer dissolved in either the dilute solution or concentrated solution is anionic, said anionic polymer may be formed by polymerisation of at least one anionic monomer alone or with other monomers. Suitable anionic monomers include ethylenically unsaturated monomers comprising carboxylic acid or sulphonic acid groups. Preferably the anionic polymer is formed from a monomer or blend of monomers comprising at least one

anionic monomer selected from the group consisting of (meth) acrylic acid, 2-acrylamido-2-methylpropane sulphonic acid, alkali metal and ammonium salts thereof.

If the polymer dissolved in either the dilute solution or concentrated solution is nonionic, said anionic polymer may be formed by polymerisation of suitable non-ionic monomers, for instance acrylamide or methacrylamide.

The polymers suitable for both the concentrated aqueous solution and dilute aqueous solution may be prepared by any convenient polymerisation process, for instance gel polymerisation, reverse phase suspension polymerisation, reverse phase emulsion polymerisation, solution polymerisation and the like. Thus suitable polymers may be provided in the form of granulated powders, beads, reverse phase emulsions, reverse phase dispersions or aqueous solutions.

The concentrated aqueous solution may be formed by dissolving any suitable water soluble polymer into water. The dilute aqueous solution of polymer may also be prepared by dissolving any suitable water soluble polymer into water or alternatively by diluting a more concentrated solution of the polymer solution.

The respective concentrated and dilute aqueous solutions may be produced therefrom by known dissolution, inversion or dilution techniques as appropriate. For instance solid particulate cationic polymer may be dissolved by dispersing the polymer particles into a flowing stream of water. Reverse phase emulsions or reverse phase dispersions of cationic polymers may be inverted into water by use of activator surfactants to form the respective aqueous solutions. Preferably the polymers dissolved in both the dilute and concentrated solutions are essentially the same polymer.

The polymers according to the invention may be prepared as substantially linear polymers or as branched or structured polymers. Structured or branched polymers are usually prepared by inclusion of polyethylenically unsaturated monomers, such as methylene-bis-acrylamide into the monomer mix, for instance as given in EP-B-

202780. Preferably however, the polymers are substantially linear and are prepared in the form of a bead or powdered product.

A particularly preferred group of polymers includes copolymers of acrylamide with at least one cationic monomer selected from the group consisting of quaternary ammonium and acid salts of dimethylaminoethyl (meth) acrylate, quaternary ammonium and acid salts of dimethylaminoethyl (meth) acrylamide and diallyldimethyl ammonium chloride, having an intrinsic viscosity of at least 4 dl/g. The cationic acrylamide polymers may comprise 10 to 90% by weight acrylamide and 10 to 90% by weight cationic monomer(s).

The aqueous composition comprising the dilute aqueous solution of cationic polymer and the concentrated solution of cationic polymer may be formed by introducing the concentrated solution of cationic polymer into a flowing stream of the dilute aqueous solution of cationic polymer. For instance in one method of preparing the aqueous composition a concentrated aqueous solution of cationic polymer is introduced directly into a conduit through which the dilute aqueous solution of cationic polymer is being conveyed towards the dosing point where the aqueous composition comprising both concentrations of polymer are metered into the suspension of solids in order to effect flocculation.

According to a preferred form of the invention wherein an aqueous suspension of suspended solids is flocculated and dewatered, an aqueous composition is introduced into the suspension. The aqueous composition comprises concentrated and dilute aqueous solutions of cationic polymer wherein the two solutions exist as discrete components of the composition. It is considered desirable that the mixture of concentrated and dilute solutions exist together as a non-homogenous composition. Therefore, in order to prevent the concentrated solution from dissipating and being diluted thus forming a homogenous solution of polymer at a single concentration, it is desirable to substantially reduce any mixing of the aqueous composition prior to being introduced into the suspension. One way that undesirable mixing of the aqueous composition can be avoided is by ensuring that

there are no mixing or pumping stages after the concentrated and dilute solutions have been combined. In addition it may further be desirable for the conduit to have a relatively smooth inner surface and the avoidance short radius bends, for example as given in pending International Application No. PCT/GB 99/00990. Another way that undesirable mixing can be avoided is to reduce the distance the aqueous composition has to travel by combining the concentrated and dilute solutions relatively close to the dosing point.

It is desirable that the aqueous composition comprising dilute and concentrated solutions does not contain substantial amounts of undissolved polymer, for instance it is preferable that less than 5%, more preferably less than 2% by weight of total polymer contained in the aqueous composition is not in solution. In many dewatering situations the most efficient use of the polymer is achieved if the amount of undissolved polymer is less than 1%, especially less than 0.5%.

The dilute solution of cationic polymer may conveniently be prepared by dilution of a more concentrated solution of the polymer. This can be achieved by adding dilution water to a flowing stream of more concentrated solution of polymer. For instance it may be desirable to pass the more concentrated solution of cationic polymer along a conduit to a dilution stage, where dilution water is introduced into the concentrated solution. In order to achieve adequate mixing of the concentrated solution with the water so that a homogenous consistent dilute solution is obtained it may be necessary to introduce a mixing stage. The mixing stage may be for instance an in-line mixing stage, such as an in-line static mixer, a pumping stage, a screening stage or some other means that can ensure adequate mixing. Preferably once thoroughly mixed the diluted solution will be substantially homogenous.

A particularly preferred aspect of the invention relates to a process of flocculating and dewatering an aqueous suspension of suspended solids by introducing into the suspension an aqueous composition comprising concentrated and dilute polymer solutions wherein the aqueous composition is formed by,



- (a) passing a concentrated solution of polymer to a dilution stage where the solution is combined with dilution water to form a dilute solution,
- (b) passing the diluted solution through a mixing stage, selected from an in-line mixer, a pumping stage and screening stage, and
- (c) introducing a concentrated solution of polymer into the dilute aqueous solution.

The concentrated polymer solution, which is diluted to form the dilute polymer solution may be drawn from the same reservoir of concentrated polymer solution which is subsequently combined with the dilute solution in forming the said aqueous composition according to the invention. There may be some mixing of the dilute and concentrated polymer solutions provided that this does not result in the aqueous composition becoming substantially homogenous.

Thus in a particularly preferred process for preparing the aqueous composition a concentrated aqueous solution of cationic polymer contained in a holding vessel is passed through a conduit to a dilution stage and subsequent mixing stage thus providing the dilute aqueous solution. Concentrated aqueous solution of cationic polymer contained in the holding vessel from said holding vessel is passed by means of a second conduit directly into the dilute aqueous solution of cationic polymer. A typical arrangement for carrying out the preparation of the aqueous composition according to this aspect of the invention is shown in Figure 1.

In Figure 1 the following key applies,

- [1] Holding vessel containing concentrated cationic polymer solution
- [2] Conduit conveying concentrated cationic polymer solution to dilution stage
- [3] Dilution water line
- [4] Pump
- [5] Conduit conveying dilute cationic polymer solution
- [6] Conduit conveying concentrated polymer solution
- [7] Aqueous composition comprising concentrated and dilute aqueous solutions of cationic polymer

- [8] Sewage sludge line
- [9] Dewatering stage
- [10] Dosing point of aqueous composition into the sludge
- [11] Dilution stage
- [12] Pump

Thus in the scheme represented in Figure 1, aqueous concentrated cationic polymer solution is held in holding vessel [1]. Concentrated polymer solution is passed along conduit [2] towards dilution stage [11] after which the aqueous polymer solution and dilution water passed through a pump [4] where they are mixed together to ensure that a consistent diluted polymer solution is formed. The dilute aqueous polymer solution is passed along conduit [5] towards the point where concentrated polymer solution is added. A second conduit [6] from holding vessel [1] conveys concentrated cationic polymer solution into the dilute polymer solution to form the aqueous composition [7] which is passed to the dosing point [10] where the mixture of concentrated and dilute cationic polymer solutions are metered into the sewage sludge line [8]. The treated sewage sludge is then passed into the dewatering stage [9].

Alternatively the concentrated polymer solution which is combined with the dilute aqueous polymer solution may be drawn from a separate reservoir of concentrated polymer solution from that which is diluted to form the dilute aqueous polymer solution. Thus in this alternative form of the invention the opportunity exists for the concentrated polymer being a different polymer from the polymer in the dilute aqueous solution. For instance it may be desirable to combine a concentrated solution of a low molecular weight cationic polymer, having an intrinsic viscosity of below 3dl/g, with a dilute solution of a high molecular weight cationic polymer, having an intrinsic viscosity of at least 4 dl/g. The low molecular weight polymer may be a coagulant, for instance the homopolymer of diallyldimethyl ammonium chloride. The high molecular weight polymer may be a bridging flocculant, for example a copolymer of acrylamide with a suitable cationic monomer, such as the quaternary ammonium salt of dimethylaminoethyl (meth)

acrylate. A typical arrangement for conducting this alternative aspect of the invention is shown in Figure 2.

In Figure 2 the following key applies,

- [1] Holding vessel containing concentrated cationic polymer solution
- [2] Conduit conveying concentrated cationic polymer solution to dilution stage
- [3] Dilution water line
- [4] Pump
- [5] Conduit conveying dilute cationic polymer solution
- [6] Second holding vessel for concentrated cationic polymer solution
- [7] Conduit conveying concentrated polymer solution
- [8] Aqueous composition comprising concentrated and dilute aqueous solutions of cationic polymer
- [9] Sewage sludge line
- [10] Dewatering stage
- [11] Dosing point of aqueous composition into the sludge
- [12] Pump
- [13] Dilution stage

Thus in the scheme represented in Figure 2, aqueous concentrated cationic polymer solution is held in holding vessel [1]. Concentrated polymer solution is passed along conduit [2] towards dilution stage [13] after which the aqueous polymer solution and dilution water are passed through a pump [4] where they are mixed together to ensure that a consistent diluted polymer solution is formed. The dilute aqueous polymer solution is passed along conduit [5] towards the point where concentrated polymer solution is added. A second conduit [7] passes concentrated aqueous cationic polymer solution from holding vessel [6] into the dilute polymer solution to form the aqueous composition [8] which is passed to the dosing point [11] where the mixture of concentrated and diluted cationic polymer solutions are metered into the sewage sludge line [9]. The treated sewage sludge is then passed into the dewatering stage [10].

The following examples serve to illustrate the invention.

[illegible]

Example 1

Aqueous solutions of a copolymer of acrylamide with dimethylaminoethyl acrylate, methyl chloride quaternary ammonium (40/60 weight/weight), intrinsic viscosity at least 10dl/g, are prepared at 0.1, 0.125 and 0.5% concentration.

Composition 1 is prepared by introducing a 0.1% solution into 0.5% solution on a 50/50 weight/ weight basis. Composition 2 is prepared in a similar manner to composition 1 by combining a 0.1% solution with 0.5% solution on a 75/25 weight/ weight basis.

200ml aliquots of Rotherham (Yorkshire, England) sewage sludge are treated with dilute polymer (0.1%) and (0.125%), concentrated polymer (0.5%) and using composition 1 and composition 2 each at various doses of cationic polymer. The treated sludge is mixed at 2000 rpm for 15 seconds. The flocculation efficiency is measured by free drainage using a 10cm diameter sieve.

The free drainage results are shown in Table 1.

Table 1

Polymer solution	5 second filtrate volume (ml) for each dose		
	137.5 mg/l	150 mg/l	162.5 mg/l
0.1%	10.5	31	55
0.125%	4	24	50
0.5%	-	27	49
Composition 1	19	41	79
Composition 2	14	32	67

The results clearly show the advantage of using the compositions comprising a mixture of concentrated and dilute solutions of the cationic polymer.

Example 2

Example 1 is repeated, except using polymer solutions 0.1%, 0.167% and 0.5% and mixed composition of 0.1% and 0.5% (50/50) and using 250ml aliquots of Rotherham sewage sludge and subjecting the treated sludge to mixing at 7000 rpm for 15 seconds. The flocculation efficiency is measured by free drainage using a 8 cm diameter sieve. For each test the volume of filtrate is measured and adjusted to allow for the volume of each aqueous polymer dose.

The adjusted free drainage results are shown in Table 2.

Table 2

Polymer solution	5 second filtrate volume (ml) for each dose				
	100 mg/l	120 mg/l	140 mg/l	160 mg/l	180 mg/l
0.1%	73	116	159	166	149
0.167%	71	114	163	174	165
0.5%	79	124	165	176	165
(50/50) mixture of 0.1% and 0.5% polymer solutions	83	166	167	166	155

The results clearly show that optimum drainage is achieved using a lower dose of the blend of 0.1% and 0.5% polymer solutions than any of the other treatments.

Example 3

Example 2 is repeated using a copolymer of acrylamide with dimethylaminoethyl acrylate, methyl chloride quaternary ammonium (80/20 weight/weight), intrinsic viscosity at least 10dl/g prepared as a reverse phase emulsion, which has been dehydrated to form a liquid dispersion product and inverted in water to form aqueous solutions of the polymers at various concentrations. These polymer solutions are tested using 500 ml aliquots of Rotherham sewage sludge which has been diluted with water (2 parts sludge to 3 parts water) and subjecting the treated sludge to mixing at 1000 rpm (low shear) for 15 seconds. The flocculation

efficiency is measured by free drainage using a 8 cm diameter sieve. The adjusted free drainage results are shown in Table 3.

Table 3

Polymer solution	5 second filtrate volume (ml) for each dose					
	30 mg/l	40 mg/l	50 mg/l	60 mg/l	70 mg/l	80 mg/l
0.1%	175	181	246	290	296	270
0.167%	121	158	246	302	308	256
0.5%	157	206	256	314	303	262
(50/50) mixture of the 0.1% and 0.5% polymer solutions	131	158	285	322	308	256

The results clearly demonstrates that the blend of polymer solutions give an increased optimum drainage by comparison to the other treatments. This is visible from the plot of these results shown in figure 3.

#### Example 4

Example 3 is repeated except the polymer has been prepared according to the teaching of EP-A-202780 by including about 20 ppm methylenebisacrylamide with the monomer resulting in a cross-linked polymer, which exhibits an ionic regain of 40%. The treatment is as described in example 3, except the treated sludge is subjected to mixing at 4,000 rpm the flocculation efficiency is measured by free drainage using an 8 cm sieve.

The free drainage results adjusted for dose volume are shown in Table 4.

Table 4

Polymer solution	5 second filtrate volume (ml) for each dose					
	90 mg/l	100 mg/l	110 mg/l	120 mg/l	130 mg/l	140 mg/l
0.1%	195	260	345	350	345	
0.167%	223	320	347	364	361	347
0.5%	221	320	369	370	382	376
(50/50) mixture of 0.1% and 0.5% polymer solutions	303	360	387	393	370	

The results clearly demonstrate that the blend of polymer solutions and separate and sequential treatment of different concentrations exhibit on the whole improved drainage by comparison to the other treatments. A plot of these results is shown in figure 4.

#### Example 5

Example 3 is repeated except the treatment comprising the mixture of 0.1% and 0.5% polymer solutions is replaced by sequential dosing of the 0.1% and 0.5% polymer solutions, wherein the 0.1% solution is added first, followed by mixing for 5 seconds at 4,000 rpm and then applying the 0.5% polymer solution, followed by further mixing for 15 seconds at 4,000 rpm and then draining through an 8cm sieve.

The free drainage results adjusted for dose volume are shown in table 5



Table 5

Polymer solution	5 second filtrate volume (ml) for each dose (Total Polymer Dose)					
	70 mg/l	80 mg/l	90 mg/l	100 mg/l	110 mg/l	120 mg/l
0.05%	80	130	210	260	300	280
0.1%	125	230	265	320	325	310
0.167%	119	226	293	320	337	324
0.2%	113	200	288	335	342	320
0.3%	108	197	275	333	352	330
0.4%	111	210	289	347	356	335
0.5%	103	172	281	340	339	328
sequential dosing of 0.1% and 0.5% polymer solutions	129	246	314	330	331	

The results clearly demonstrate that effective dewatering of the sludge can be achieved using lower total polymer dose by applying dilute and concentrated polymer solutions by comparison to the other treatments employing single concentration polymer solutions. Thus the mixed concentration dosing enables more efficient dosing of the polymer.

#### Example 6

Aqueous solutions of a copolymer of acrylamide with dimethylaminoethyl acrylate, methyl chloride quaternary ammonium (75/25 weight/weight), intrinsic viscosity at least 10dl/g, are prepared at 0.1, 0.125 and 0.5% concentration. A mixture of 0.1% and 0.5% solution as a weight ratio of 75:25 is also prepared.

Dewatering of a de-inked paper mill sludge (0.91% solids) was evaluated using the polymer solutions at various doses. For each test the polymer was dosed to 600ml of sludge, followed by stirring for 15 seconds at 2000 rpm using a 4 blade

stirrer. The flocculation efficiency was measured using free drainage through an 8 cm sieve recording filtrate volume after 5 seconds. The free drainage results, adjusted to take into account the dose volumes are shown in table 6.

Table 6

Polymer solution	5 second filtrate volume (ml) for each dose		
	1 Kg/t	2 Kg/t	4 Kg/t
0.1%	125.5	139	108
0.125%	145.6	141.2	82.4
0.5%	148.9	147.8	115.6
75:25 mixture of 0.1% solution and 0.5% solution	145.6	161.2	102.4

As can be seen by the results of this test the mixture of dilute and concentrated polymers solutions provides improved optimum free drainage.

#### Example 7

Example 6 was repeated except instead of measuring free drainage, the treated sludge was transferred to a piston press. Pressures of 20, 40, 60 and 80 pounds per square inch (psi) were applied in 2 minute increments.

The cake produced was then weighed wet and dry to calculate the cake solids.

The results are shown in table 7.

Table 7

Polymer solution	Cake Solids %		
	1 Kg/t	2 Kg/t	4 Kg/t
0.1%	25.15	26.94	30.83
0.125%	31.82	29.84	33.09
0.5%	42.93	26.56	31.24
75:25 mixture of 0.1% solution and 0.5% solution	26.34	32.31	32.95

The results from examples 6 and 7 show that the mixed dilute and concentrated polymer solutions provide the best overall combination of free drainage and cake solids.

#### Example 8

A suspension of China Clay is prepared and used 4% (weight/volume) in 2g/l sodium chloride solution. The tests are carried out on 500 ml aliquots of the China Clay suspension and mixed with various doses of polymer solutions of specified concentrations at 500 rpm impeller speed. The duration of mixing is for 15 seconds for single doses and simultaneous doses.

The flocculated China Clay is for each test transferred to a 500 ml measuring cylinder immediately upon completing the mixing stage. The time taken for the solid liquid interface (mud line) to pass between the 3 cm and 8 cm level is measured. A sedimentation rate in cm/minute is calculated and shown for each total polymer dose in table 8.

In each test the polymer is a copolymer of acrylamide with sodium acrylate, with a monomer ratio by weight of 70:30.

Table 8

Polymer solution	Sedimentation rate (cm/min)		
	3 mg/l	4 mg/l	5 mg/l
0.05%	21	40.7	57.1
0.0833%	15	41.2	55.6
0.25%	14.8	36.5	43.8
50:50 mixture of 0.05% solution and 0.25% solution	27.2	55.9	83.6

The results show that the mixed concentrated and dilute polymer solutions provide the best sedimentation rates. This is clearly shown in figure 5.

#### Example 9

Example 8 is repeated comparing two stage addition of dilute and concentrated polymer solutions with simultaneous addition, two stage addition of two dilute solutions and single stage addition of a dilute solution.

The duration of mixing is and for 15 seconds for single doses and simultaneous doses and for the two stage dosing of concentrated and dilute polymer solutions, the first dose is applied followed by mixing for 5 seconds followed by the second dose and then mixing for a further 15 seconds.

A sedimentation rate in cm/minute is calculated and shown for each total polymer dose in table 9.

Table 9

Polymer solution	Sedimentation rate (cm/min)			
	3 mg/l	4 mg/l	5 mg/l	6 mg/l
0.05%	10.2	14.6	20.7	38
0.05% two stage addition	12.7	20	29.2	42.1
0.05%:0.25% (50:50) two stage addition	20.2	33.4	38.5	47.1
50:50 mixture of 0.05% solution and 0.25% solution	15.7	32.1	43.7	45.2

The results show that the mixed concentrated and dilute polymer solutions and two stage addition of dilute and concentrated polymer solutions out perform the single dose of dilute polymer solution or the two stage dose of dilute polymer solution. This is clearly visible from the plots shown in figure 6.

### Claims

1. A process of flocculating and dewatering an aqueous suspension of suspended solids comprising, introducing into the suspension,

- (a) a concentrated polymer solution and,
- (b) a dilute polymer solution,

characterised in that the concentrated and dilute polymer solutions are introduced into the substrate substantially simultaneously.

2. A process according to claim 1 in which the concentrated polymer solution and dilute polymer solution are introduced into the suspension as an aqueous composition comprising,

- (a) a dilute aqueous solution of polymer and,
- (b) a concentrated solution of polymer,

wherein the dilute solution and concentrated solution exist as substantially discrete components.

3. A process according to claim 2 in which the aqueous composition comprises,

- (a) 25 to 99%, by weight of the dilute aqueous solution of polymer and,
- (b) 1 to 75% by weight of the concentrated solution of polymer.

4. A process according to claim 2 in which the aqueous composition comprises,

- (a) 40 to 99%, by weight of the dilute aqueous solution of polymer and,
- (b) 1 to 60% by weight of the concentrated solution of polymer.

5. A process according to any of claims 1 to 4 in which the dilute aqueous solution has a concentration of polymer of below 0.3% by weight.

6. A process according to any one of claims 1 to 5 in which the dilute aqueous polymer solution comprises a cationic polymer, an anionic polymer or a nonionic polymer.

7. A process according to any of claims 1 to 6 in which the concentrated aqueous solution has a concentration of polymer of between 0.4 and 1.0%, by weight.

8. A process according to any one of claims 1 to 7 in which the concentrated aqueous solution comprises a cationic polymer, an anionic polymer or a nonionic polymer.

9. A process according to any one of claims 1 to 8 in which the polymer dissolved in the concentrated solution is either co-ionic with the polymer dissolved in the dilute solution or non-ionic.

10. A process according to any one of claims 1 to 9 in which polymer dissolved in either the dilute solution or the concentrated solution is cationic and has been formed from a monomer or blend of monomers comprising at least one cationic monomer selected from the group consisting of quaternary ammonium and acid salts of dimethylaminoethyl (meth) acrylate, quaternary ammonium and acid salts of dimethylaminoethyl (meth) acrylamide and diallyldimethyl ammonium chloride.

11. A process according to any one of claims 1 to 10 in which the polymer(s) dissolved in either the dilute solution or the concentrated solution is anionic and has been formed from a monomer or blend of monomers comprising at least one anionic monomer selected from the group consisting of (meth) acrylic acid, 2-acrylamido-2-methylpropane sulphonic acid, alkali metal and ammonium salts thereof.

12. A process according to any one of claims 1 to 11 in which the polymer(s) dissolved in either the dilute solution or the concentrated solution is nonionic and has been formed from acrylamide or methacrylamide.

13. A process according to any one of claims 1 to 12 in which the cationic polymer dissolved in each of the dilute and concentrated aqueous solutions is a copolymer of acrylamide and at least one cationic monomer selected from the group consisting of quaternary ammonium and acid salts of dimethylaminoethyl (meth) acrylate, quaternary ammonium and acid salts of dimethylaminoethyl (meth) acrylamide and diallyldimethyl ammonium chloride, having an intrinsic viscosity of at least 4 dl/g.

14. A process according to any one of claims 2 to 13 in which the aqueous composition comprising the dilute aqueous solution of polymer and the concentrated solution of polymer is formed by introducing the concentrated solution of polymer into a flowing stream of the dilute aqueous solution of polymer.

15. A process according to claim 14 in which the dilute aqueous solution of polymer is formed by diluting a flowing stream of the concentrated aqueous solution of polymer with dilution water.

16. A process according to any one of claims 2 to 15 in which the aqueous composition is formed by,

(a) passing a concentrated solution of polymer to a dilution where the solution is combined with dilution water to form a dilute solution,

(b) passing the diluted solution through a mixing stage, selected from pumping and screening stages, and

(c) introducing a concentrated solution of polymer into the dilute aqueous solution.

17. A process according to claim 16 in which the concentrated polymer solution, which is diluted to form the dilute polymer solution in step (a) is drawn from the same reservoir of concentrated polymer solution introduced into the dilute solution in step (c).

18. A process according to claim 16 in which the concentrated solution of polymer in step (a) is drawn from a different reservoir of concentrated polymer solution introduced into the dilute solution in step (c).

19. A process according to any one of claims 1 to 18 in which the dewatering process is selected from the group consisting of dewatering sewage sludge, dewatering a mineral suspension, dewatering a paper mill sludge, dewatering a deinked cellulosic sludge and a papermaking process.



Figure 1

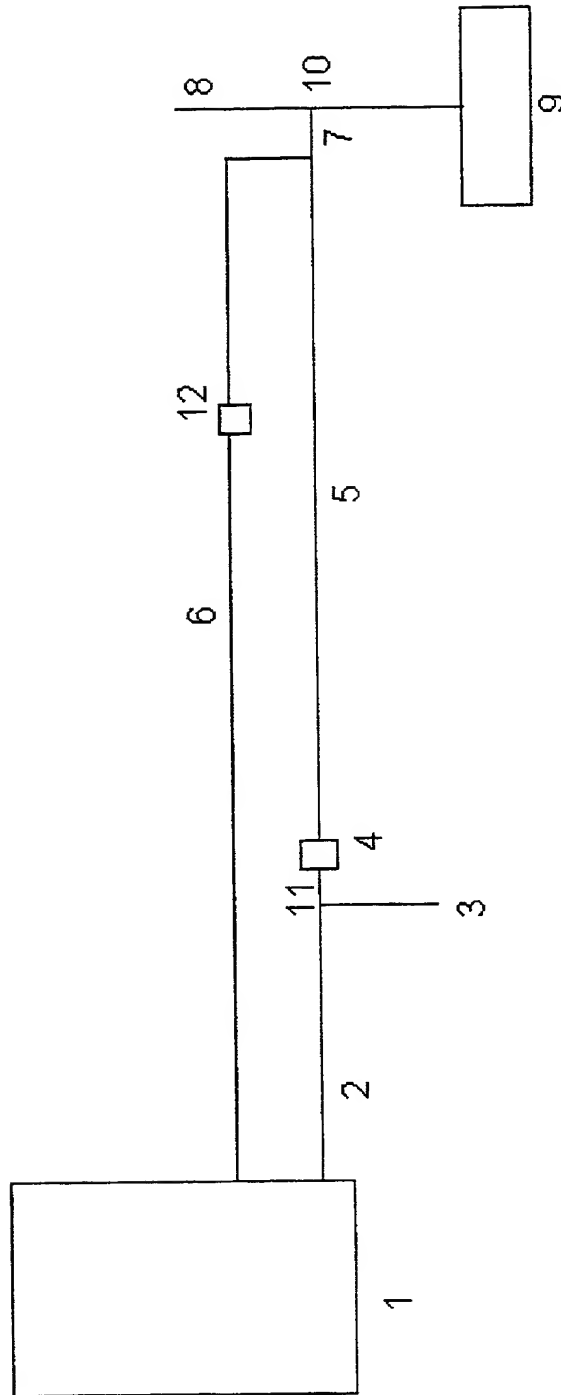


Figure 2

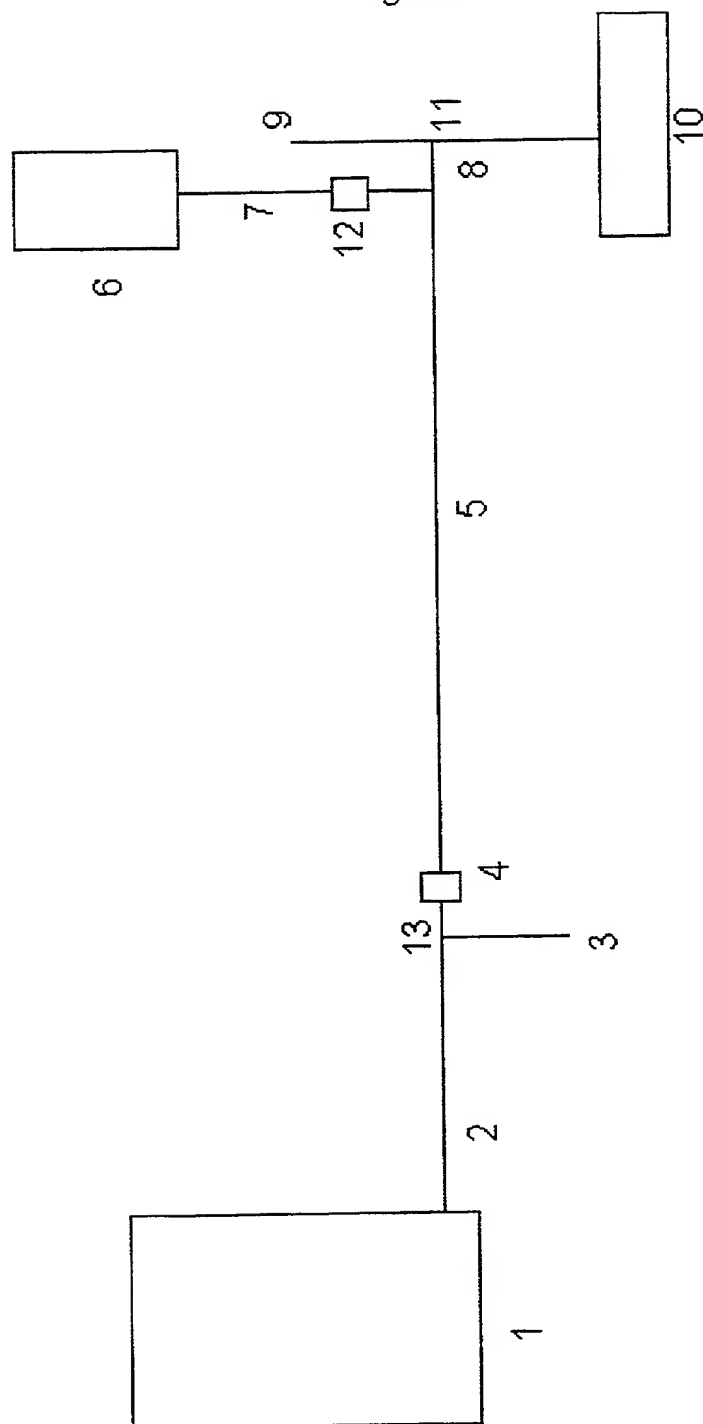
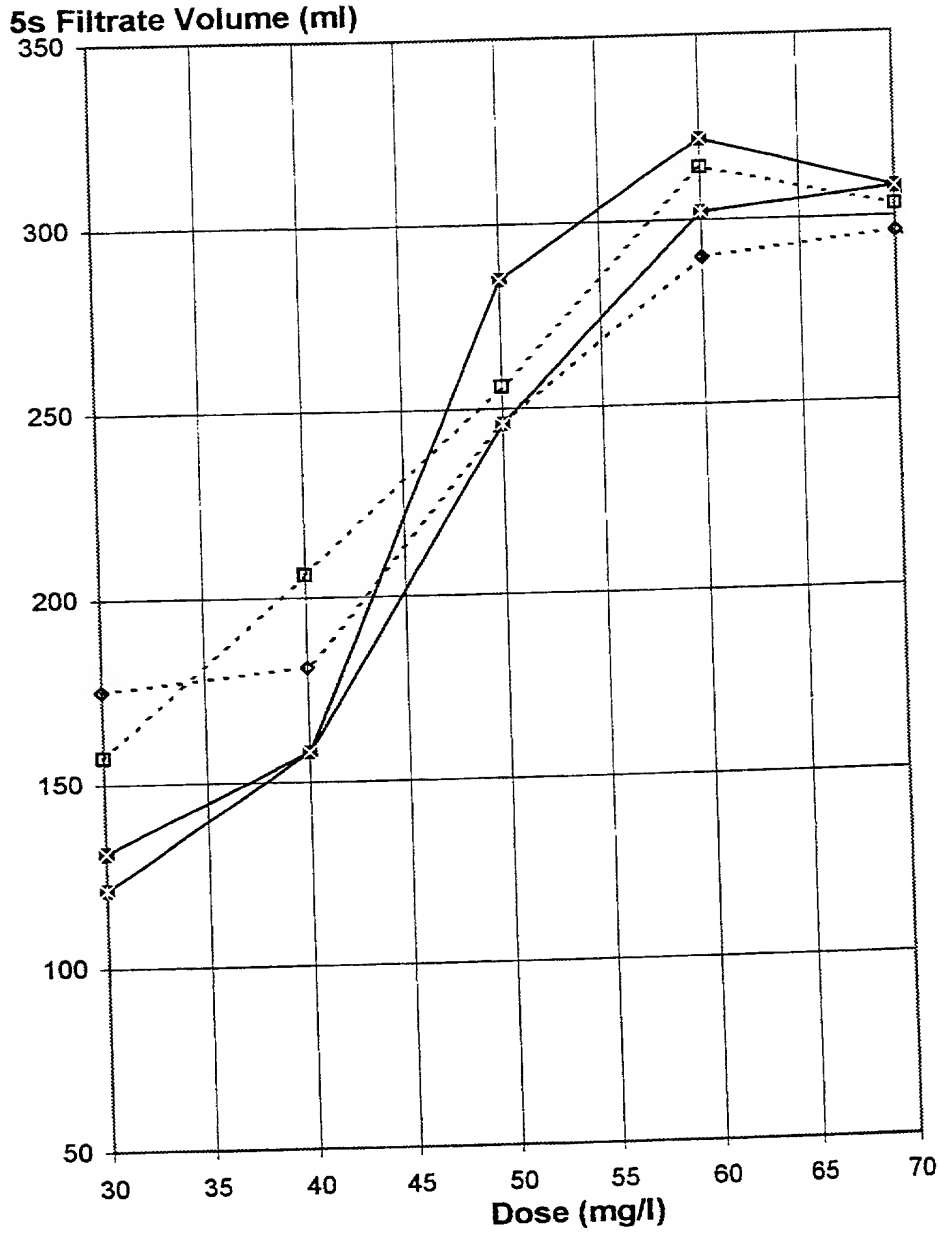


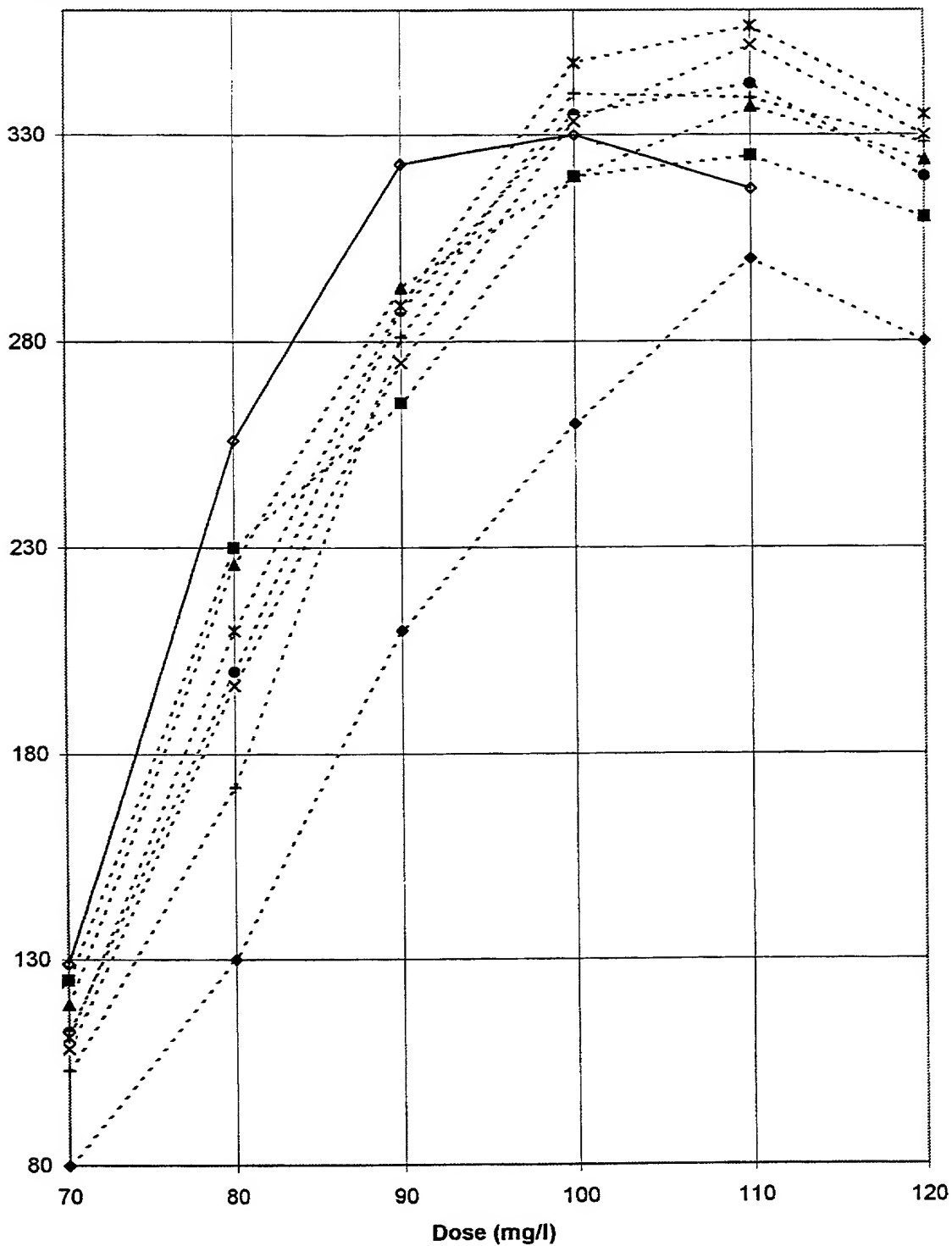
Figure 3



---◇--- 0.10%      ---□--- 0.50%      ---×--- 0.167%      ---×--- 0.1 + 0.5%

Figure 4

5s filtrate vol. (ml)



---◆---0.05%	---■---0.1%	---▲---0.167%
---●---0.2%	---×---0.3%	---*---0.4%
---+---0.5%	—◇—0.1% then 0.5%	

Figure 5

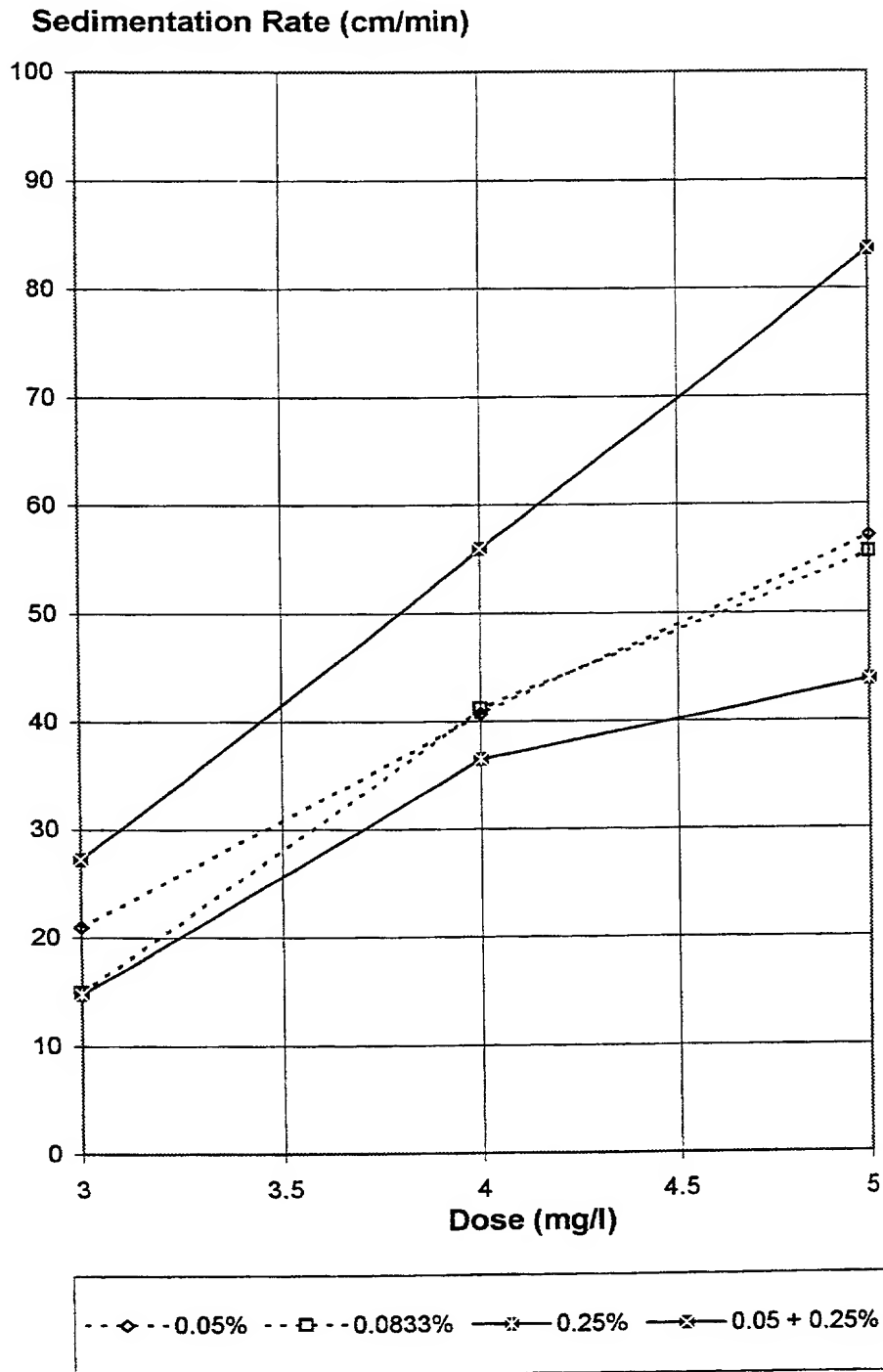
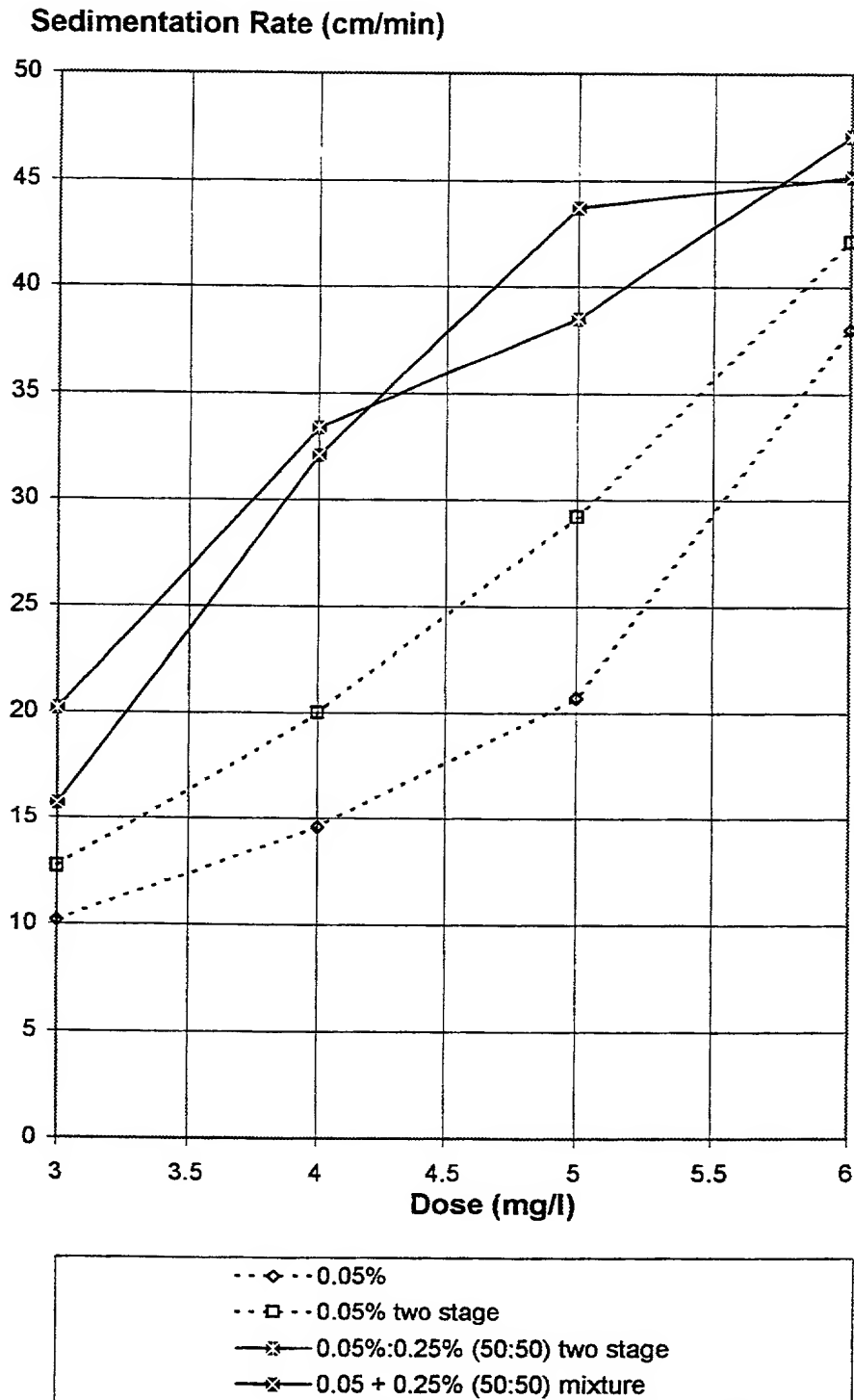


Figure 6



**DECLARATION AND POWER OF ATTORNEY FOR U.S. PATENT APPLICATIONS**

☒ Original      ☐ Supplemental      ☐ Substitute      ☐ PCT

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if more than one name is listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled

**Process For Flocculating Suspensions**

which is described and claimed in:

- ☐ the attached specification.
- ☐ the specification in U.S. Application No. \_\_\_\_\_, and as amended on \_\_\_\_\_ (if applicable).  
(day/month/year) (day/month/year)
- ☒ the specification in International Application No. **PCT/EP00/06292**  
filed **05/07/2000**  
(day/month/year)
- assigned U.S. Application No. \_\_\_\_\_ (if applicable), and as amended
- ☐ under PCT Article 19 on \_\_\_\_\_ (if applicable)  
(day/month/year)
- ☐ under PCT Article 34 on \_\_\_\_\_ (if applicable)  
(day/month/year)
- ☐ and further amended on \_\_\_\_\_ (if applicable)  
(day/month/year)

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose all information which is known by me to be material to the patentability of this application as defined in 37 C.F.R. § 1.56.

I hereby claim foreign priority benefits under 35 U.S.C. § 119 (a)-(d) of any foreign application(s) for patent or inventor's certificate or of any PCT international application(s) designating at least one country other than the United States of America listed below and have also identified below any foreign application(s) for patent or inventor's certificate or any PCT international application(s) designating at least one country other than the United States of America relating to this subject matter having a filing date before that of the application on which priority is claimed:

COUNTRY/REGION (OR PCT)	APPLICATION No.	FILING DATE (day/month/year)	PRIORITY CLAIMED	
<b>Great Britain</b>	<b>9916748.8</b>	<b>19/07/1999</b>	<input checked="" type="checkbox"/> Yes	<input type="checkbox"/> No
			<input type="checkbox"/> Yes	<input type="checkbox"/> No
			<input type="checkbox"/> Yes	<input type="checkbox"/> No
			<input type="checkbox"/> Yes	<input type="checkbox"/> No
			<input type="checkbox"/> Yes	<input type="checkbox"/> No

I hereby claim the benefit under 35 U.S.C. § 119 (e) of any United States provisional application(s) listed below:

APPLICATION NO.	FILING DATE (day/month/year)
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I hereby claim the benefit under 35 U.S.C. § 120 of any United States application(s) or PCT international application(s) designating the United States listed below and, insofar as the application discloses and claims subject matter in addition to that disclosed in the prior copending application, I acknowledge the duty to disclose all information known by me to be material to patentability as defined in 37 C.F.R. § 1.56 which became available between the filing date of the prior application and the national or PCT international filing date of this application:

U.S. APPLICATION No.	FILING DATE (day/month/year)	STATUS		
		<input type="checkbox"/> Patented	<input type="checkbox"/> Pending	<input type="checkbox"/> Abandoned
		<input type="checkbox"/> Patented	<input type="checkbox"/> Pending	<input type="checkbox"/> Abandoned
		<input type="checkbox"/> Patented	<input type="checkbox"/> Pending	<input type="checkbox"/> Abandoned
		<input type="checkbox"/> Patented	<input type="checkbox"/> Pending	<input type="checkbox"/> Abandoned
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			<input type="checkbox"/> Patented
			<input type="checkbox"/> Pending
			<input type="checkbox"/> Abandoned



I hereby appoint the following attorneys and agents, associated with Customer No. 000324, each of them with full power of substitution, revocation and appointment of associates, to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith:

Luther A. R. Hall (Reg. No. 27,337), JoAnn L. Villamizar (Reg. No. 30,598), Kevin T. Mansfield (Reg. No. 31,635), Jacob M. Levine (Reg. No. 32,509), David R. Crichton (Reg. No. 37,300) and Michele A. Kovalski (Reg. No. 37,865).

Address all correspondence associated with Customer No. 000324 to ***Ciba Specialty Chemicals Corporation, Patent Department, 540 White Plains Road, P.O. Box 2005, Tarrytown, NY 10591-9005.***

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C. § 1001, and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full name of sole  
or first joint inventor     **Steven WEIR**

Inventor's signature \_\_\_\_\_ Date \_\_\_\_\_  
(day/month/year)

Residence     29 St Thomas Gardens,  
Bradley,  
Huddersfield,  
West Yorkshire,  
HD2 1SL,  
GB

Citizenship     **British**

Post Office Address     Same as above

2-00

Full name of second  
joint inventor, if any Mark John STRINGER

Inventor's signature M.J. Stringer Date 9/12/01  
(day/month/year)

Residence 15 Woodhall Croft,  
Pudsey,  
Leeds,  
West Yorkshire,  
LS28 7TU,  
GB GBX

Citizenship British

Post Office Address same as above

Full name of third  
joint inventor, if any

Inventor's signature \_\_\_\_\_ Date \_\_\_\_\_  
(day/month/year)

Residence

Citizenship

Post Office Address same as above

20060709 090000

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Full name of sole  
or first joint inventor Steven WEIR

Inventor's signature Steven Weir Date 12/11/2001  
(day/month/year)

Residence 29 St Thomas Gardens,  
Bradley,  
Huddersfield,  
West Yorkshire,  
HD2 1SL,  
GB

GBX

Citizenship **British**

Post Office Address Same as above

Full name of second  
joint inventor, if any

**Mark John STRINGER**

Inventor's signature

Date

(day/month/year)

Residence

**15 Woodhall Croft,  
Pudsey,  
Leeds,  
West Yorkshire,  
LS28 7TU,  
GB**

Citizenship

**British**

Post Office Address

same as above

Full name of third  
joint inventor, if any

Inventor's signature

Date

(day/month/year)

Residence

Citizenship

Post Office Address

same as above